Docket No. UF-375 Serial No. 10/602,394

## In the Claims

This listing of claims will replace all prior versions and listings of claims in this application.

I (currently amended). A peptide that is biologically active at melanocortin receptors comprising an AGRP(109-118) analogue template of SEQ ID NO:3, [[and]] wherein AGRP (111-113) residues of the AGRP (109-118) template are substituted with a melanocortin agonist-based bioactive determinant sequence sequences which have been substituted for the analogous template sequences; wherein

a) the melanocortin agonist-based bioactive determinant sequence is selected from the group consisting of:

- i) Tip-Arg-Phe;
- ii) Trp-Arg-DPhe;
- iii) Phe-Arg-Trp;
- iv) DPhc-Arg-Trp;
- v) His-Phe-Arg-Trp; and
- vi) His-DPhc-Arg-Trp.

2 (original). The peptide according to claim 1, wherein the peptide is of any SEQ ID NOS:4-7, 9, and 10.

3 (currently amended). The peptide according to claim 1, wherein at least one of the amino acids in the melanocortin agonist-based bioactive determinant sequence includes at least one amino acid-substituted within the sequence is substituted with an amino acid selected from the group consisting of Ala; Atc. Bip; Lys; Nal(1'); Nal(2'); (pl)Phe; and Tic.

4 (cancelled).

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Docket No. UF-375 Serial No. 10/602,394

5 (currently amended). The peptide according to claim [[4]]3, wherein the peptide is of any SEQ ID NOS:24-43.

6 (currently amended). The peptide according to claim 1, wherein the peptide further comprises a lactam bridge [[which]] that is substituted for the disulfide bridge of the AGRP(109-118) analogue template.

7 (original). The peptide according to claim 6, wherein the peptide is of any SEQ ID NOS:2 and 11.

8 (original). The peptide according to claim 6, wherein the peptide further comprises a second and a third bioactive determinant sequences at the N-terminal and C-terminal, respectively, wherein the second bioactive determinant sequence at the N-terminal is Ser-Tyr-Ser-Nle amino acid residues and the third bioactive determinant sequence at the C-terminal is Lys-Pro-Val amino acid residues.

9 (cancelled).

10 (cancelled).

H (cancelled).

12 (cancelled).

13 (currently amended). A pharmaccutical composition comprising a peptide that is biologically active at melanocortin receptors comprising an AGRP(109-118) analogue template of SEO ID NO:3, wherein AGRP (111-113) residues of the AGRP (109-118) template are substituted with a [[and]] melanocortin agonist-based bioactive determinant sequence sequences which-have

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Docket No. UF-375 Serial No. 10/602,394

been substituted for the analogous template sequences, and a pharmaceutically acceptable carrier or diluent, wherein

- a) the melanocortin agonist-based bioactive determinant sequence is selected from the group consisting of:
  - Trp-Arg-Phc;
  - ii) Trp-Arg-DPhe;
  - iii) Phe-Arg-Trp;
  - iv) DPhc-Arg-Trp;
  - v) His-Phe-Arg-Trp; and
  - vi) His-DPhe-Arg-Trp.

14 (original). The pharmaceutical composition according to claim 13, wherein the peptide is of any SEQ ID NOS:4-7, 9, and 10.

15 (currently amended). The pharmaceutical composition according to claim 13, wherein at least one of the residues of the melanocortin agonist-based bioactive determinant sequence includes at least one-amino-acid substituted-within the sequence is substituted with an amino acid selected from the group consisting of Ala; Atc; Bip; Lys; Nal(1'); Nal(2'); (pl)Phe; and Tig.

16 (cancelled).

17 (original). The pharmaceutical composition according to claim 16, wherein the peptide is of any SEQ ID NOS:24-43.

18 (currently amended). The pharmaceutical composition according to claim 13, wherein the peptide further comprises a lactain bridge [[which]] that is substituted for the disulfide bridge of the AGRP(109-118) template.

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Docket No. UF-375 Serial No. 10/602,394

19 (original). The pharmaceutical composition according to claim 18, wherein the peptide is of any SEQ 1D NOS:2 and 11.

20 (original). The pharmaceutical composition according to claim 18, wherein the peptide further comprises a second and a third bioactive determinant sequences at the N-terminal and C-terminal, respectively, wherein the second bioactive determinant sequence at the N-terminal is Ser-Tyr-Ser-Nie amino acid residues and the third bioactive determinant sequence at the C-terminal is Lys-Pro-Val amino acid residues.

- 21 (cancelled).
- 22 (cancelled).
- 23 (cancelled).
- 24 (cancelled).
- 25 (cancelled).
- 26 (cancelled).
- 27 (cancelled).

28 (previously presented). The composition of claim 13, wherein the composition is an oral composition.

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